Applicants: Stoffel et al. Serial No.: 10/824,644 Filed: April 13, 2004

Amendment and Response to April 17, 2007 Office Action

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## **AMENDMENTS TO THE SPECIFICATION**

## Please replace the paragraph beginning at page 3, line 26 with the following:

Figure 2: Predicted precursor structure and tissue expression of mouse miR-375. (A) RNA secondary structure prediction was performed using Mfold version 3.1 (SEQ. ID. NO. 31). The miRNA sequence is underlined. There is complete homology between mouse and human sequences. (B) Tissue expression of miR-375 and -376. Total RNA (30 μg) were isolated from mouse tissues for Northern blots and probed for the indicated miRNA. (C) Northern blots of total RNA (10μg) isolated from purified pancreatic islets, MIN6 cells and total pancreas. High expression levels were detected in mouse pancreatic islets. A tRNA probe was used as a loading control.

## Please replace the paragraph beginning at page 4, line 24 with the following:

Figure 5: The miR-375 target site in the 3'UTR of Mtpn is responsible for inhibition of gene expression by miR-375 (SEQ. ID. No. 1). (A) Sequence of the target site in the 3'UTR of myotrophin inserted within the Renilla luciferase 3' UTR. The mutant construct (Mtpn-MUT) (SEQ. ID. NO. 70) is identical to the WT construct (Mtpn-WT) (SEQ. ID. NO. 69) except for five point mutations (bold) disrupting base-pairing at the 5' end of miR-375.

(B) MIN6 cells were transiently transfected with either reporter construct in addition to 2'-O-methyl-oligoribonucleotides complementary to miR-375 (2'-O-methyl-375) or a control 2'-O-oligoribonucleotide (2'-O-methyl-GFP).